EVALUATION OF ANTINOCICEPTIVE EFFECT OF THE KETOROLAC-TOPIRAMATE COMBINATION IN THE RAT FORMALIN TEST

Claudia Cervantes-Durán¹, Erandi Sánchez-Serrano², Marcia Yvette Gauthereau-Torres² and Luis Fernando Ortega-Varela³*

¹Bachelor of Information Technology in Sciences, National School of Higher Studies, Morelia Unit, National Autonomous University of Mexico, Morelia, Michoacan
²Division of Graduate Studies, Faculty of Medical and Biological Sciences “Dr. Ignacio Chávez”, Michoacan University of San Nicolás de Hidalgo, Morelia, Michoacan
³Faculty of Public Health and Nursing, Michoacan University of San Nicolás de Hidalgo, Morelia, Michoacan

Combination therapy approaches to manage acute and chronic pain are commonly used. To characterize the interaction between ketorolac and topiramate in the formalin test, female Wistar rats (200-300 g) were submitted to 1% formalin test. Antinociceptive effect was determined by the administration of ketorolac (3, 10, 30 and 100 mg/kg), topiramate (25, 50, 100 and 200 mg/kg) and their combination by oral route; or ketorolac (25, 50, 100 and 200 µg/paw), topiramate (10, 30, 100 and 300 µg/paw) and their combination by local peripheral route. Isobolographic analysis was used in a fixed dose combination (0.5:0.5) to analyze the nature of the interaction of the combination based on the ED₅₀ of ketorolac (62.8±21.9 mg/kg) and topiramate (24.3±8.5 mg/kg) by oral administration; or the ED₃₀ of ketorolac (67.0±7.97 µg/paw) and topiramate (391.3±51.7 µg/paw) by local peripheral route. Combination of these two drugs significantly reduced the number of flinches in second phase of the test. Theoretical ED₅₀ of the oral combination (ED₅₀T) was 43.5±11.7 mg/kg. Experimentally, the ED₅₀ of the combination (ED₅₀E) had a significantly lower value: 16.7±4.0 mg/kg; indicating the presence of supra additive effects (interaction index was 0.38). For local peripheral route, ED₃₀T was 229.21±64.99 µg/kg. Experimentally, the ED₃₀E had a significantly lower value: 62.00±15.72 µg/paw; indicating synergistic effects (interaction index was 0.27). Results show that oral and local peripheral administration of the combination can interact synergistically to reduce inflammatory pain in the rat formalin test.

INTRODUCTION

Nonsteroidal anti-inflammatory drugs (NSAIDs) are a class of therapeutic agents widely used to treat pain, inflammation and fever¹. Ketorolac is a NSAID which exhibits a potent analgesic activity in the treatment of moderate to severe pain. Ketorolac is non-narcotic but provides opioid-level pain management, thereby reducing narcotic requirements². Experimental observations show that ketorolac exhibits a potency similar to those of indomethacin and diclofenac in inhibiting cyclooxygenase 1 and cyclooxygenase 2 besides prostaglandin synthesis. The activation of the NO–cyclic GMP–ATP-sensitive K⁺ channel pathway plays an important role in ketorolac antinociception³. Ketorolac may be prescribed to control cancer-associated pain and is used as an analgesic during and after cancer surgeries². However, like other NSAIDs, ketorolac has been associated with significant gastrointestinal, renal, and cardiovascular risks⁴.

*Corresponding author: Luis Fernando Ortega-Varela, E-mail: lfortega@umich.mx
In the search for alternatives that solve the demand for effective pain treatment, a large number of preclinical and clinical investigations have been carried out to find analgesic substances or analgesic combinations that increase their potency, reducing the risk of undesirable effects. One of the possible ways for improving inflammatory pain treatment is to use alternative analgesics (drugs from various pharmacologic groups with primary indications other than pain that have shown to be effective in certain pain states). Antiepileptic drugs (AEDs) are among the most important alternative analgesics. In this sense, carbamazepine, gabapentin and topiramate, have been used for neuropathic pain treatment due to their neuromodulatory effect on pain perception. Moreover, recent findings suggest that topiramate applied systemically and/or locally could be useful as an analgesic against inflammatory pain. Previous reports of combinations with ketorolac or topiramate show that it is feasible to increase the analgesic effects in inflammatory pain models and the use of both drugs could be a rational target for combination. The purpose of this study was to assess the possible synergistic interaction between ketorolac and topiramate orally or peripherally administrated in the rat formalin test.

**MATERIALS AND METHODS**

**Animals**
Experiments were performed on female Wistar rats (200-300 g). Rats were kept under controlled conditions of temperature (22±2°C) and light (12 h:12 h) with ad libitum access to water and food. All experiments followed the guidelines on ethical standards for investigation of experimental pain in animals and Mexican regulation. The institutional ethics committee approved all experiments (CB/2018/V-220). Rats were euthanized in a CO₂ chamber at the end of the experiment.

**Drugs**
Ketorolac ((±)-5-Benzoyl-2,3-dihydro-1H-pyrrolizine-1-carboxylic acid) and topiramate (2,3:4,5-Bis-O-(1-methylethylidene)-36-D-fructo-pyranose sulfamate) were purchased from Sigma (St. Louis, Mo., USA). Both drugs were dissolved in saline.

**Measurement of antinociceptive activity**
Nociception was assessed using the formalin test. Rats were placed in open observation chambers (Plexiglas) for acclimatization (30 min); then they were gently restrained for the injection of 50 µl of diluted formalin (1%) into the dorsal surface of the right hind paw with a 30-gauge needle. The animals were returned to the chambers immediately after formalin injection and nociceptive behavior was registered. Mirrors were placed in each chamber to enable unhindered observation. Nociceptive behavior was quantified as the number of flinches of the injected paw during 1-min periods every 5 min, up to 60 min after injection. Flinching is a simple and reliable parameter of pain behavior and one producing high scores and was characterized as rapid and brief withdrawal, or as flexing of the injected paw. Formalin-induced flinching behavior was biphasic. The phase 1 (acute phase, 0–10 min) was followed by a relatively short inactive period, which was then followed by the phase 2 (prolonged tonic response, 15-60 min). Rats were euthanized in a CO₂ chamber at the end of the experiment.

**Study design**
For the systemic study, animals received a unique dose of vehicle or increasing concentrations of either ketorolac (25, 50, 100 and 200 mg/kg), topiramate (10, 30, 100 and 300 mg/kg) or the ketorolac-topiramate combinations (Table 1) orally, 10 min before (50 µl) of subcutaneous formalin injection. For the local peripheral study, rats received a single subcutaneous administration of vehicle or increasing doses of either ketorolac (25, 50, 100 and 200 µg/paw), topiramate (10, 30, 100 and 300 µg/paw) or the ketorolac-topiramate combination (as indicated in Table 1) 10 min before formalin injection. For all routes of administration, doses and time of administration were selected on the basis of previous studies in this model. Rats in all groups were observed regarding behavioral or motor function changes induced by the treatments. This was assessed, but not quantified, by testing the animals’ ability to stand and walk in a normal posture, as proposed elsewhere.
Table 1: Maximum effect of oral and local peripheral administration of ketorolac and topiramate, alone or in combination in phase 1 in the formalin test.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>% antinociception maximum effect in phase 1 oral route</th>
<th>% antinociception maximum effect in phase 1 local-peripheral route</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ketorolac (200 mg/kg; 200 µg/paw)</td>
<td>57.33 ± 2.73</td>
<td>24.93 ± 12.06</td>
</tr>
<tr>
<td>Topiramate (300 mg/kg; 300 µg/paw)</td>
<td>77.07 ± 4.17</td>
<td>71.28 ± 2.34</td>
</tr>
<tr>
<td>Ketorolac-topiramate combination (43.92 mg/kg; 7.08 µg/paw)</td>
<td>55.76 ± 5.72</td>
<td>59.45 ± 2.55</td>
</tr>
</tbody>
</table>

Data analysis

All results are presented as mean ± S.E.M. of six animals per group. Time-courses of antinociceptive responses of individual drugs and their combinations were constructed by plotting the mean number of flinches as a function of time. To compare three or more experimental groups, we used one-way analysis of variance (ANOVA), followed by a Student Newman-Keuls test. The area under the curve (AUC) of the number of flinches against time, for each case, was calculated by the trapezoidal method. Dose-response data are presented as percent of antinociception, calculated from AUC of phase 2 of the formalin test; according to the following equation:

\[
\text{% Antinociception} = \frac{\text{AUC}_{\text{vehicle}} - \text{AUC}_{\text{drugs}}}{\text{AUC}_{\text{vehicle}}} \times 100
\]

The dose-response curves were constructed and the experimental points fitted using least-square linear regression. ED\textsubscript{50} ± standard error (S.E.M.) was calculated according to the method described by Tallarida\textsuperscript{7}. It has been previously demonstrated that, for evaluation of the interaction between analgesic drugs, isobolographic analysis is a critical tool\textsuperscript{18,22}. In the present study, we used this method to determine the nature of drug interactions between ketorolac and topiramate when administered orally and local peripherally. In this analysis we assumed that the combination of drugs represents equieffective doses of the individual drugs. Thus, from the dose-response curves of each individual agent, the dose resulting in 50% of the effect (ED\textsubscript{50} value) can be determined. However, considering a maximal effect of 100% as the total suppression of formalin-induced flinches, it appeared that topiramate was unable to achieve a 50% response in the local peripheral route, and thus a calculation of ED\textsubscript{50} value was not possible for this case. Therefore, we estimated the ED\textsubscript{30} value instead of the ED\textsubscript{50} value only for local peripheral combination.

To determine if the oral interaction between both drugs given in combination was synergistic, additive, or antagonistic, the theoretical additive ED\textsubscript{50} value was estimated from the log dose-response curve of each compound administered individually, considering that the observed combined effect results from the addition of the individual effects of each component. This theoretical ED\textsubscript{50} value of the combination was then compared with the experimental ED\textsubscript{50} values to determine if there is a significant difference\textsuperscript{18,22}. The experimental ED\textsubscript{50} value was obtained by oral coadministration of fractions of the ED\textsubscript{50} respective value for each model: 1/2, 1/4, 1/8, 1/16. Isobolographic analysis was then used to characterize the antinociceptive interaction between ketorolac and topiramate in the formalin test. The theoretical and experimental ED\textsubscript{50} values of the studied combinations were also compared by calculating the interaction index (\(\gamma\)) as follows:\textsuperscript{20}:

\[
\gamma = \frac{\text{ED}_{50} \text{ value of combination (experimental)}}{\text{ED}_{50} \text{ value of the combination (theoretical)}}
\]
The interaction index indicates the portion of ED<sub>50</sub> value of the individual compound that accounts for the corresponding ED<sub>50</sub> value in the combination, that is, values near to unity correspond to an additive interaction, values higher than 1 indicate an antagonistic interaction, and values lower than 1 indicate a synergistic interaction<sup>23</sup>. For the determination of the local peripheral interaction, we used the same procedure but using de ED<sub>50</sub> values instead of ED<sub>50</sub>.

RESULTS AND DISCUSSION

Results
Oral and peripheral antinociceptive effect of ketorolac and topiramate
Administration of ketorolac or topiramate significantly reduced formalin-induced nociceptive behavior by oral or local peripheral routes during phases 1 and 2 of the formalin test. However, since drugs did not produce a dose-dependent reduction of the nociceptive behavior in phase 1 (data not shown), only phase 2 was submitted to further analysis. Maximum percent of antinociception in phase 1 are summarized in table 1. Figure 1 shows the typical time course of the formalin test and the effects of topiramate and ketorolac administered separately orally (A) and peripherally (B). In the dose-response curve for ketorolac, doses of 3, 10, 30 and 100 mg/kg were orally administered. Phase 2 of the test showed a maximum antinociceptive effect for oral ketorolac of about 51% and the effects observed were dose-dependent (Fig. 2A). For topiramate, doses of 25, 50, 100 and 200 mg/kg were administered orally (Fig. 2B). In phase 2, the effect of topiramate was dose-dependent with an efficacy of about 61%.

Fig. 1: Temporary courses of the formalin test obtained by oral administration (po) (A) and local peripheral (lp) route (B). The data are the mean of 6 rats ± SEM.

Fig. 2: Dose-response curves in the second phase of formalin test for ketorolac (Ket 3, 10, 30 and 100 mg/kg) and topiramate (Top 25, 50, 100 and 200 mg/kg) administered orally (po). In panel A, the figure shows the results for ketorolac. In panel B, the effects of topiramate are observed. The results were dose-dependent and the maximum effect recorded was 51.2% for ketorolac and 64.6% for topiramate (n= 6; *p< 0.05 versus V= Vehicle, for one-way ANOVA followed of Student-Newman-Keuls).
Fig. 3: Dose-response curves in the second phase of the formalin test for ketorolac (Ket 25, 50, 100 and 200 µg/paw) and topiramate (Top 10, 30, 100 and 300 µg/paw) administered by local peripheral route (lp). The figure shows in panel A, the results for ketorolac. In panel B, the effects of topiramate are observed. The results were dose-dependent and the maximum effect registered was 55.7% for ketorolac and 30.0% for topiramate (n = 6; *p < 0.05 versus V= Vehicle, for one-way ANOVA followed of Student-Newman-Keuls.

Combination effects

Co-administration of ketorolac and topiramate induced a dose-dependent increase in antinociception percent in phase 2 after oral and local peripheral administration (Fig. 4A). In the second phase of the test, the oral theoretical effective dose 50 (ED₅₀T) of the combination was 43.5±11.7 mg/kg, while the ED₅₀E was 16.7±4.0 mg/kg, which is significantly lower than the theoretical one, showing that the interaction is synergistic (Fig. 4B).

The dose-response curve of the local peripheral combination was carried out in phase 2 of the test using the doses described in table 2. The highest dose (229.2 µg/paw) corresponds to the theoretical effective dose 30 (ED₃₀T). Once the doses of the combination were established by the calculations of the isobolographic analysis, they were subjected experimentally to the formalin test. In the second phase, the observed effect was dose-dependent. The dose corresponding to the ED₃₀T had an antinociceptive effect of about 57.4% as an experimental result (Fig. 5A).

When the data obtained from the dose-response curve of the combination of topiramate and ketorolac administered by the local peripheral route in phase 2 of the test were compared with the previously calculated theoretical values, the ED₃₀T, whose value is 229.21 µg/paw, was different from the ED₃₀E (62.31±15.72 µg/paw), the experimental values were lower than those expected for a purely additive interaction (Fig. 5B).

Table 2: Effect of local peripheral and oral administration of ketorolac and topiramate alone or in combination in the formalin test.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Oral administration ED₅₀, mg/kg</th>
<th>Local Peripheral administration ED₃₀, µg/paw</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ketorolac</td>
<td>62.84 ± 21.96</td>
<td>67.08 ± 7.97</td>
</tr>
<tr>
<td>Topiramate</td>
<td>24.30 ± 8.52</td>
<td>391.34 ± 129.74</td>
</tr>
<tr>
<td>Theoretical combination</td>
<td>43.57 ± 11.77</td>
<td>229.21 ± 64.99</td>
</tr>
<tr>
<td>Experimental combination</td>
<td>16.70 ± 4.01*</td>
<td>62.31 ± 15.72*</td>
</tr>
<tr>
<td>Interaction index</td>
<td>0.38</td>
<td>0.27</td>
</tr>
</tbody>
</table>

ED₅₀: effective dose resulting in a 50% reduction of control response. ED₃₀: effective dose resulting in a 30% reduction of control response. Data are the mean ± SEM of the estimate. *Significantly different from the theoretical combination data (p < 0.05), by the Student’s t-test.
Fig. 4: In panel A, the dose-response curve of the oral combination (po) in the second phase of the formalin test is shown. The ED$_{50}$ corresponding to 43.57±11.77 mg/kg produced an antinociceptive effect of 72.9% in phase 2 of formalin test ($n=6$; *$p<0.05$ versus V= Vehicle, by one-way ANOVA followed by Student-Newman-Keuls). In panel B, the isobologram of the oral (po) interaction of ketorolac and topiramate is shown. Horizontal and vertical bars indicate SEM. The oblique line between the x- and y-axis is the theoretical additive line. The point in the middle of this line, indicated by T, is the theoretical additive point calculated from the individual drug ED$_{50}$ values. The experimental point, indicated by E, is the ED$_{50}$ actually observed with the combination. The experimental ED$_{50}$ point lies far below the additive line, indicating a significant synergistic interaction, ($p<0.05$) as determined by the Student’s t-test.

Fig. 5: In panel A, the dose-response curve of the local peripheral combination (lp) in the second phase of the formalin test is shown. The ED$_{30}$ corresponding to a total of 222.20±64.99 µg/paw produced an antinociceptive effect of 57.4% in phase 2 of formalin test ($n=6$; *$p<0.05$ versus V= Vehicle, by one-way ANOVA followed by Student-Newman-Keuls). In panel B, the isobologram of the local peripheral (lp) interaction of ketorolac and topiramate is shown. Horizontal and vertical bars indicate SEM. The oblique line between the x- and y-axis is the theoretical additive line. The point in the middle of this line, indicated by T, is the theoretical additive point calculated from the individual drug ED$_{30}$ values. The experimental point, indicated by E, is the ED$_{30}$ actually observed with the combination. The experimental ED$_{30}$ point lies far below the additive line, indicating a significant synergistic interaction. The experimental ED$_{30}$ point lies far below the additive line, indicating a significant synergistic interaction, ($p<0.05$) as determined by the Student’s t-test.
For each ketorolac-topiramate combination, the interaction index ($\gamma$) was lower than 1 (0.38 and 0.27 for oral and peripheral combinations, respectively). Analysis of the interaction index (Table 2) showed an increase in potency after oral and local peripheral administration, almost threefold for every combination employed, highlighting their synergistic effects.

Discussion

Antinociceptive effect of ketorolac

In the present study, the administration of oral or local peripheral ketorolac significantly decreased the number of flinches in the formalin test, as it has been thoroughly demonstrated in previous studies with ketorolac\textsuperscript{1,12,25}. In the clinic, ketorolac is the most potent NSAID used to treat any pain associated with inflammation, especially postoperative pain, renal colic, arthritis, lumbago, headache and pain from cancer. It is prescribed for use in the short term and for the management of severe acute pain that requires immediate analgesia\textsuperscript{26}. However, long-term use of any NSAID, including ketorolac, may be associated with peptic ulcer and some other systemic side effects such as coagulation disorder, nephrotoxicity, as well as severe disability in cardiac, cerebral or hepatic functions\textsuperscript{25,26}.

Ketorolac is a bioavailable drug for several routes of administration such as oral and parenteral, for both routes it has a significant analgesic potency, as demonstrated by the present data. Analgesic efficacy of ketorolac has been studied extensively for the treatment of moderate to severe pain. Its oral administration provides analgesia that is equal to or better than aspirin or acetaminophen\textsuperscript{29}. Unlike other NSAIDs, ketorolac is a potent analgesic with excellent aqueous solubility; it does not irritate the tissues. It is useful for its administration by different conventional routes: iv, im, oral, topical and rectal, but at the same time opens the way for new therapeutic possibilities\textsuperscript{28}.

However, ketorolac mainly exerts its effects through the inhibition of COX (1 and 2 isoforms), with a higher affinity for COX-1. The inhibition of COX decreases the production of prostaglandins, thromboxane and prostacyclin from arachidonic acid. Prostaglandins are involved in the nociceptive pathway by sensitizing the afferent nerves\textsuperscript{29}. Since it is not selective for any of the COX isoforms, it inhibits both the formation of pro-inflammatory PGs in peripheral nociceptors (anti-inflammation and analgesia: desired pharmacological effects) and the production of "protective" PGs from COX-1. This phenomenon explains many of the adverse effects secondary to the administration of ketorolac\textsuperscript{30}.

In this study, the combination of ketorolac and topiramate required only 30% or less of the individual dose of ketorolac, which means that in that proportion its adverse effects could be reduced and the time of use of ketorolac could be extended. In this scenario, the combination could be useful for the treatment of chronic diseases that induce pain and inflammation.

Antinociceptive effect of topiramate

Antiepileptic drugs, like topiramate, are widely used in neuropathic pain treatment, but there is substantial preclinical evidence about their efficacy against inflammatory pain\textsuperscript{6,8}. Our study agrees with previous reports, as we observed that either systemic or local peripheral administration of topiramate produced a dose-related antinociception in rats submitted to noxious stimulation with formalin in the second phase of the test\textsuperscript{9,12}.

Topiramate has several pharmacological properties that may contribute to its antinociceptive effect that include: inhibition of voltage activated Na\textsuperscript{+} channels and some L-type high-voltage activated Ca\textsuperscript{2+} channels; a negative modulatory effect on the AMPA/kainate subtypes of GluRs; a positive effect on some GABA\textsubscript{A} receptors; and inhibition of the carbonic anhydrase isoenzymes CA-II and CA-IV\textsuperscript{31}.

Antinociceptive effect of topiramate and ketorolac combination

Combinations of analgesic drugs are often prescribed with the intent of enhancing the therapeutic effect without increasing, or possibly reducing the side effects\textsuperscript{22}. The main objective of this study was to determine the type of pharmacological interaction between ketorolac and topiramate administered orally and peripherally. According to our understanding, this is the first study about the
interaction of these drugs. Oral and local peripheral administration of the ketorolac-topiramate combination significantly reduced the nociceptive behavior induced by formalin. It was also observed that the antinociceptive response produced by the administered drugs is dose-dependent, at least in phase 2 of the test. All these results agree with previous investigations that show that the systemic administration of NSAIDs like ketorolac and antiepileptics, such as topiramate, produce a dose-dependent antinociceptive activity in several animal models. In this sense, when administered locally, both drugs were effective against inflammatory and neuropathic pain, and some systemic side effects could be avoided. The interaction of these drugs has not been previously assessed, however, ketorolac has been tested in analgesic combinations with morphine, tizanidine, B vitamins, and tramadol; while topiramate has been tested with diacerhein, gabapentin and tramadol. The interaction index of this combination, a measure of the degree of synergism, produced similar level by oral and local peripheral routes (0.38 and 0.27, respectively); showing a threefold rise in the potency of the combinations when compared with the theoretical effective doses, indicating possible advantages by both administration routes in the management of pain at different conditions.

The mechanism of this antinociceptive interaction remains to be elucidated, nevertheless, if we consider our findings and previous reports, we can suggest that the sites and mechanism of action of both drugs seem to be complementary. This combination involves, at least, the inhibition of COX 1 and 2 by ketorolac, which prevents the synthesis of prostaglandins involved in peripheral and central sensitization. On the other hand, topiramate inhibits neuronal excitability by blocking sodium and calcium channels, promoting GABAergic activity. Thus, the combination of topiramate and ketorolac could diminish, at least by two mechanisms of action, the transmission of nociception.

No side effects were observed during the present experiments, suggesting that this combination has a favorable side-effect profile. Commonly reported adverse effects of topiramate are dizziness, ataxia, and disorientation. However, previous data of our group indicates that topiramate did not alter rota-rod performance at doses employed here. In fact, Shank et al. reported the same effect of this drug even in doses higher than 2000 mg/kg. In addition, the reduction of doses in the synergistic/additive combinations could reduce dose-dependent, drug-specific adverse effects and improve treatment tolerability.

**Conclusion**

Our results show that the ketorolac-topiramate combination produces a functional synergic activity to reduce inflammatory pain at different levels of pain transmission. The reduction of almost two thirds of the dose requirements with no exacerbation of side effects profile suggests that this combination could be useful against inflammatory pain states.

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تقييم التأثير المضاد للالتهاب لمزيج كيتورولاك - توبيراميت في اختبار الفورمالين بالجرذان

كلوديا سيرفاتيس دوران1 - إرانيدي سانشيز سيرانو2 - مارسيا إييفيت غاوثريو توريس3
لويس فرناندو أورتيجا-فاريلا2

1بكالوريوس تكنولوجيا المعلومات ، المدرسة الوطنية للدراسات العليا ، وحدة موريلبا ، جامعة المكسيك الوطنية المستقلة. موريلبا ، ميشواكان
2شعبة الدراسات العليا بكلية العلوم الطبية والبيولوجية "D. إنجينيوس شافير". جامعة ميشواكان ، سان نيكولاس دي هيدالغو. موريلبا ، ميشواكان

كلية الصحة العامة والتمريض ، جامعة ميشواكان داي سان نيكولاس دي هيدالغو. موريلبا ، ميشواكان

يشير استخدام أساليب العلاج المركب لإدارة الألم الحاد والمزمن. لتصنيف التفاعل بين كيتورولاك و توبيراميت في اختبار الفورمالين ، تم إجراء اثنان فحص و يستغرق 300 يوم عد الجفّل (200-300 جم) لاختبار الفورمالين بنسبة 2%. تم تحديد التأثير المضاد للمسببات عن طريق إعطاء كيتورولاك (30, 20, 10، و 50 مجم/ كجم) ، توبيراميت (25, 50، 100، و 200 ميكروجرام/ للمخلب) ، توبيراميت (10، 20، 30 و 40 ميكروجرام/ للمخلب) وتوليفهم عن طريق الفم؛ أو كيتورولاك (25، 50، 100، و 200 ميكروجرام/ للمخلب) و توبيراميت (3، 4، 5 و 6 ميكروجرام/ للمخلب) وتوليفهم عن طريق الفم. تم استخدام تحليل Isobolographic في جرعات ثابتة للمزيج (0، 5، 10 مجم/ كجم) للتعرف على التفاعل أو التداخل بناءً على ED50 من كيتورولاك (8، 11، 14، 17، 20 مجم/ كجم) و توبيراميت (3، 4، 5، 6 مجم/ كجم) عن طريق الفم؛ أو من كيتورولاك (27، 30، 33، 36 مجم/ كجم) و توبيراميت (1، 2، 3، 4 مجم/ كجم) عن طريق الفم. 

بكلية الصحة العامة والتمريض، جامعة ميشواكان داي سان نيكولاس دي هيدالغو. موريلبا ، ميشواكان

نظرًاً للتوازن الممتاز في عدد الجفّل في المرحلة الثانية من الاختبار، كان ED50 للنواة الناجمة (ED50E) للنواة الناجمة ED50 (7 3.5±11.7 مجم/ كجم) ومجموعة ED50 للنواة الناجمة ED50 (7 3.5±11.7 مجم/ كجم) . يشير إلى وجود تأثيرات فوق مساحة (مؤشر التفاعل) 0.38

بالنسبة للنواة الناجمة ED30 T (7 29.2±11.7 مجم/ كجم). من النواة الناجمة ED30 T (7 29.2±11.7 مجم/ كجم) و توبيراميت (7 4، 5، 6 مجم/ كجم) . يشير إلى تأثيرات تازية (كان مؤشر التفاعل 0.27). تظهر النتائج أن إعطاء الفموي والموضعي للمزيج يمكن أن يتفاعل بشكل تازير في تقليل الألم الالتهابي في اختبار الفورمالين للفئران.